HIGH YIELD REVIEW POINTS FOR REVISION BY DR ABDUL BASIT ZUBAIR Ear

- 1. **Conductive hearing loss** shows bone conduction better than air conduction (Rinne negative) and lateralizes to the affected ear in Weber test.
- 2. **Sensorineural hearing loss** has air conduction better than bone (Rinne positive) but Weber lateralizes to the unaffected ear.
- 3. **Otosclerosis** is a hereditary condition of abnormal bone remodeling at the stapes footplate, often affecting young women; hearing may paradoxically improve in noisy environments.
- 4. **Chronic suppurative otitis media (CSOM)** may cause persistent ear discharge and hearing loss; unsafe CSOM may form cholesteatoma leading to erosion of nearby structures.
- 5. **Cholesteatoma** is an epithelial cyst in the middle ear or mastoid with keratin debris; it can erode ossicles and cause complications like facial palsy or brain abscess.
- 6. **Glomus tumors** are vascular tumors (paragangliomas) that can cause pulsatile tinnitus and hearing loss; they may invade cranial nerves IX-XII.
- 7. **Eustachian tube dysfunction** leads to middle ear negative pressure and recurrent otitis media, particularly in children due to anatomical predisposition.
- 8. **Facial nerve injury** can occur during mastoid or middle ear surgery; the tympanic segment is especially at risk.
- 9. **Perilymph fistula** presents with sudden sensorineural hearing loss and vertigo after trauma or straining, due to leakage of inner ear fluid.
- 10. **Tympanometry**: Type A is normal; type B (flat curve) suggests middle ear effusion; type C suggests negative middle ear pressure.

Vertigo, Tinnitus, Otalgia

- 11. **BPPV** is caused by dislodged otoliths in semicircular canals and is diagnosed by Dix-Hallpike maneuver, which reproduces vertigo and nystagmus.
- 12. **Ménière's disease** involves endolymphatic hydrops, causing episodes of vertigo, tinnitus, and sensorineural hearing loss, often with ear fullness.
- 13. **Vestibular neuronitis** is a self-limiting post-viral inflammation of the vestibular nerve causing vertigo without hearing loss.
- 14. **Tinnitus** may be pulsatile (suggesting vascular cause like AV malformation or glomus tumor) or non-pulsatile (sensorineural origin).
- 15. **Otitis media** can cause referred otalgia due to shared innervation with the glossopharyngeal and vagus nerves.

- 16. **Ramsay Hunt syndrome** (herpes zoster oticus) causes painful vesicles on ear and facial palsy; CN VII is affected.
- 17. **Labyrinthitis** includes both vestibular and cochlear symptoms following infection; it can follow viral URTI.
- 18. **Superior semicircular canal dehiscence** causes vertigo with loud sounds (Tullio phenomenon); confirmed via CT.
- 19. **Tinnitus** without hearing loss might require imaging to exclude retrocochlear pathology like acoustic neuroma.
- 20. **TMJ dysfunction or tonsillitis** can refer pain to the ear, causing otalgia with normal otoscopic findings.

Nose

- 21. **Nasal polyps** are edematous mucosa, often bilateral and associated with asthma, chronic rhinosinusitis, and aspirin sensitivity (Samter's triad).
- 22. **Allergic rhinitis** presents with sneezing, nasal congestion, and clear discharge; often with allergic shiners and nasal crease.
- 23. **Septal hematoma** after nasal trauma must be drained to prevent cartilage necrosis and resultant saddle nose deformity.
- 24. **Woodruff's plexus** is a venous plexus on the posterior nasal floor; bleeding here is usually profuse and difficult to control.
- 25. **Hereditary hemorrhagic telangiectasia** causes recurrent epistaxis due to fragile blood vessels; look for mucocutaneous telangiectasias.
- 26. **Allergic fungal sinusitis** presents with thick allergic mucin and opacified sinuses with hyperdensities on CT.
- 27. **Rhinosporidiosis** is a chronic infection caused by *Rhinosporidium seeberi*, presenting as a red, friable nasal mass that bleeds on touch.
- 28. **Unilateral nasal discharge** in a child is a foreign body until proven otherwise, especially if foul-smelling.
- 29. **Kiesselbach's plexus (Little's area)** is the common site for anterior epistaxis, especially in children and dry climates.
- 30.**Sphenopalatine artery** is often the culprit in posterior epistaxis and may require endoscopic ligation.

Throat

- 31. **Peritonsillar abscess** presents with fever, trismus, and uvula deviation away from the affected side; requires drainage.
- 32. **Retropharyngeal abscess** in children presents with neck stiffness, fever, and dysphagia; may compress the airway.
- 33. **Laryngomalacia** causes inspiratory stridor in infants due to floppy supraglottic structures; often resolves by 2 years.
- 34. **Epiglottitis** presents with high fever, drooling, and respiratory distress; "thumb sign" on lateral neck X-ray.
- 35. **Vocal nodules** occur due to voice abuse; bilateral and symmetric on vocal cords, vs. polyps which are unilateral.
- 36. **Reinke's edema** is swelling of the vocal cords in smokers, causing husky voice; common in middle-aged women.
- 37. **Recurrent laryngeal nerve palsy** causes hoarseness and a fixed vocal cord; most often iatrogenic during thyroid surgery.
- 38. **Arytenoid dislocation** from intubation trauma causes persistent hoarseness and breathy voice.
- 39. **Laryngeal papillomatosis** is due to HPV types 6 and 11; causes multiple vocal fold growths and hoarseness in children.
- 40. **Supraglottic cancers** metastasize early due to rich lymphatics; glottic cancers present early due to voice change.

Dysphagia

- 41. **Zenker's diverticulum** is a pharyngoesophageal pouch leading to dysphagia, regurgitation, and halitosis.
- 42. **Plummer-Vinson syndrome** occurs in iron deficiency anemia with esophageal webs and glossitis; pre-malignant.
- 43. **Schatzki ring** causes intermittent dysphagia to solids; associated with hiatal hernia.
- 44. Achalasia causes failure of LES relaxation; barium swallow shows bird-beak appearance.
- 45. **Cricopharyngeal spasm** presents with intermittent globus sensation; often functional.

Neck

46. **Thyroglossal cyst** moves upward with tongue protrusion; common midline neck mass in children.

- 47. **Branchial cleft cyst** is a lateral neck mass along anterior border of SCM; arises from incomplete obliteration of branchial clefts.
- 48. **Reactive cervical lymphadenopathy** is the most common cause of neck mass in children; often secondary to infection.
- 49. **Virchow's node** in the supraclavicular region may indicate gastric or abdominal malignancy.
- 50. **Midline neck swellings** in children are usually thyroglossal duct cysts; surgical removal requires Sistrunk procedure.

OPHTHALMOLOGY

- 1. Thyroid Eye Disease (Graves' Orbitopathy)
 - **Presentation:** Bilateral proptosis, lid retraction, restrictive extraocular movements (inferior & medial rectus most common), NO fever/redness.
 - **Key Feature: Lid lag, exophthalmos**, may have optic neuropathy.
 - **Diagnosis:** Clinical + **TSI antibodies** (Thyroid Stimulating Immunoglobulin).
 - Management: Mild → lubricants, steroids; Severe → orbital decompression.

2. Orbital vs. Preseptal Cellulitis

Feature Orbital Cellulitis	Preseptal Cellulitis
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Location	Post-septal (deep)	Pre-septal (superficial)
Proptosis	Yes	No
Eye Movement s	Restricted & painful	Normal
Vision	May be affected	Normal
Cause	Sinusitis (most common), trauma	Skin infection (stye, bug bite)
Treatment	IV antibiotics (Vancomycin + Ceftriaxone), may need surgery	Oral antibiotics

3. Dacryocystitis (Lacrimal Sac Infection)

- Acute: Painful medial canthal swelling, pus regurgitation on pressure.
- **Chronic:** Constant tearing (**epiphora**), recurrent infections.
- Newborns: Nasolacrimal duct obstruction → Crigler massage (90% resolve by 1 year).
- Treatment:
 - O Acute → IV antibiotics (Augmentin).
 O Chronic → Dacryocystorhinostomy (DCR).

4. Eyelid Lesions

Lesion	Features	Treatment
Hordeolum (Stye)	Painful, lid margin , staph infection	Warm compresses, drainage
Chalazion	Painless, non-marginal , granuloma	Steroid injection, excision
Blepharitis	Crusting, itching, chronic	Lid hygiene, erythromycin

5. Conjunctivitis

Туре	Features	Treatment
Bacterial	Purulent discharge, sticky lids	Fluoroquinolones (Ciprofloxacin)
Viral (Adeno)	Watery discharge, preauricular LN	Supportive, highly contagious
Allergic (Vernal)	Itching, ropy discharge , papillae	Mast cell stabilizers (Opatanol)

6. Corneal Pathology

• Herpetic Keratitis: Dendritic ulcer (fluorescein stain), photophobia, treat with acyclovir.

- Pseudomonas Keratitis (Contact Lens Users): Ring infiltrate, rapid progression → Ciprofloxacin drops.
- Fungal Keratitis (Farmers): Feathery margins, trauma with vegetative matter →
 Natamycin.

7. Uveitis

- Symptoms: Pain, photophobia, ciliary congestion, miosis, low IOP.
- Key Sign: Keratic precipitates (KPs) + cells in anterior chamber.
- Treatment: Steroids (Prednisolone drops) + Cycloplegics (Atropine).

8. Glaucoma

Туре	Features	Treatment
Open- Angle	Painless, gradual vision loss, cupped disc	Prostaglandins (Latanoprost) → βblockers (Timolol)
Angle- Closure	Sudden pain, nausea, fixed middilated pupil, high IOP	IV Mannitol → Pilocarpine → Laser iridotomy

9. Retinal Disorders

- Diabetic Retinopathy:
 - o **NPDR:** Microaneurysms, hemorrhages, hard exudates.
 - o PDR: Neovascularization \rightarrow Pan-retinal photocoagulation (PRP).
- Retinal Vein Occlusion (CRVO): Flame hemorrhages, venous dilatation → Macular edema → Anti-VEGF (Ranibizumab).
- Retinal Detachment: Flashes & floaters → curtain-like vision loss → Urgent surgery.

10. Neuro-Ophthalmology

• Optic Neuritis: Pain on eye movement, RAPD (Marcus Gunn pupil), normal fundus initially → MRI (MS risk).

Horner's Syndrome: Ptosis + Miosis + Anhidrosis → Sympathetic chain lesion (Pancoast tumor, carotid dissection).

CN Palsies:

CN III (Oculomotor): Ptosis, down & out eye, pupil involved (aneurysm).
 CN IV (Trochlear): Vertical diplopia, head tilt (trauma).

o CN VI (Abducens): Cannot abduct (raised ICP, diabetes).

11. Trauma & Emergencies

- Chemical Burns: Immediate irrigation (30 mins), pH check.
- Ruptured Globe: Do NOT patch, shield → Urgent surgery.
- Corneal FB: Remove with needle, check for rust ring.

12. Pediatric Ophthalmology

- Congenital Nasolacrimal Duct Obstruction: Crigler massage, resolves by 1 year.
- Amblyopia: Occlusion therapy (patch good eye).
- Strabismus: Hirschberg test, esotropia (inward), exotropia (outward).

13. Vitamin A Deficiency

- Early Sign: Conjunctival xerosis.
- Late Signs: Bitot's spots, corneal xerosis → keratomalacia (blindness).
- Treatment: High-dose Vitamin A.

14. Drug Mechanisms

- Atropine: Cycloplegic + Mydriatic (contraindicated in glaucoma).
- Pilocarpine: Miotic (opens angle in glaucoma).
- Latanoprost: ↑ Uveoscleral outflow (PG analogue).
- Fluoroquinolones (Ciprofloxacin): Inhibit DNA gyrase.

Key Mnemonics:

- CRVO: "Blood & thunder" retina (hemorrhages).
- CRAO: Cherry-red spot (central retinal artery occlusion).
- Papilledema: Bilateral disc swelling (ICP个, headache).

PHARMACOLOGY



Respiratory

- Acute bronchial asthma Salbutamol
- **COPD** Anticholinergics (Tiotropium, Ipratropium)
- Aspirin-induced asthma Leukotriene inhibitors (Zafirlukast, Montelukast)

Infections

- MRSA Vancomycin
- VRSA Linezolid
- Amebiasis / Giardiasis / Trichomoniasis / C. difficile Metronidazole
- Syphilis Benzathine Penicillin
- Systemic fungal infection Amphotericin-B
- HSV/Shingles Acyclovir
- Malaria in pregnancy Chloroquine

CNS

Absence seizure – Ethosuximide
 GTCS – Valproate
 Partial seizures – Carbamazepine

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Eclampsia – Magnesium sulfate Status epilepticus – Lorazepam

• Anaphylaxis – Epinephrine

Endocrine/Metabolic

- **DM Type 1** Insulin
- **DM Type 2** Metformin
- **DKA** Crystalline insulin
- **Hyperprolactinemia** Bromocriptine
- **Hypothyroidism** Levothyroxine
- **Hyperthyroidism in pregnancy** Propylthiouracil

CVS

- **AF** Digoxin
- **SVT** Adenosine
- Prinzmetal angina Nitroglycerin
- Pulmonary edema Furosemide
- HTN in pregnancy Methyldopa, Labetalol

Others

- Migraine Sumatriptan
- Gout (acute) NSAIDs
- Cerebral edema Mannitol
- Osteoporosis prevention Bisphosphonates
- Motion sickness Scopolamine
- Traveler's diarrhea Diphenoxylate, Norfloxacin

Antidotes

- Paracetamol N-acetylcysteine
- Benzodiazepine Flumazenil
- Heparin Protamine sulfate
 Warfarin Vitamin K (delayed), FFP (immediate)
 Opioids Naloxone
- Organophosphates Atropine > Pralidoxime
- **Digoxin** Digoxin Fab
- Magnesium sulfate Calcium gluconate
- **Isoniazid** Pyridoxine (B6)

⚠ Common Drug Side Effects

- ACE inhibitors Cough, angioedema, hyperkalemia
- Amphotericin B Nephrotoxicity
- Atropine Dry mouth, constipation
- Ciprofloxacin Tendon rupture
- Clozapine Agranulocytosis
- Lithium Hypothyroidism, nephrogenic DI
- INH Peripheral neuropathy, hepatitis
- **Vincristine** Peripheral neuropathy
- Methotrexate Pancytopenia, mucositis
- Amiodarone QT prolongation, hypothyroidism, phototoxicity

A Mechanism of Action

- Aminoglycosides / Tetracyclines Inhibit 30S
- Macrolides / Clindamycin / Chloramphenicol Inhibit 50S
- Vancomycin Inhibits peptidoglycan synthesis

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- **Sulfonamides** Inhibit folate synthesis
- **Methotrexate** Inhibits dihydrofolate reductase
- Rifampin Inhibits RNA polymerase
- INH Inhibits mycolic acid synthesis
- Fluoroquinolones Inhibit DNA gyrase (topoisomerase)
- **Statins** HMG CoA reductase inhibitors

Allopurinol – Xanthine oxidase inhibitor

Theophylline – PDE inhibitor $\rightarrow \uparrow$ cAMP

Steroids - Inhibit phospholipase A2

Finasteride – 5α -reductase inhibitor

ANATOMY

Here is a systematic summary of the high-yield ANATOMY points from the NLE Essence PDF:

One Neuroanatomy

- CSF production:
- → 20 mL/hr; total daily production ≈ 450 mL
- → Formed by **choroid plexus**
- Sensory Receptors:
- → Pacinian corpuscles Vibration, deep pressure
- → **Ruffini corpuscles** Steady pressure, warmth
- → End bulb of Krause Cold

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ightarrow Muscle spindle – Stretch reflex ightarrow

Golgi tendon organ – Tendon reflex

- CNS Lesions & Structures:
- → Chorea Basal ganglia lesion
- → Intention tremor Cerebellum lesion

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- → **Resting tremor** Substantia nigra (e.g., Parkinson's)
- → Accommodation center Cerebral cortex
- → Corneal reflex Pons
- → Pupillary light reflex Midbrain
- → Wernicke-Broca connection Arcuate fasciculus
 - Hemorrhages on CT Scan:
- → **Subdural hematoma** Crescent-shaped
- → Epidural hematoma Biconvex/lens-shaped
- → Subarachnoid hemorrhage Severe headache + blood-tinged CSF

♦ General Anatomy & Bone

- Most common site of aortic aneurysm Infrarenal aorta
- Rupture of AAA Left retroperitoneum is most common CT angiography Investigation of choice for AAA
- Exostosis of EAC (Surfer's ear):
- → Benign bony growth, multiple, bilateral, sessile, due to cold water exposure
- → CT shows broad-based lesion
 - Osteoma:
- → Unilateral, pedunculated, outer part of EAC
- → Associated with **Gardner syndrome**

A Microscopic Anatomy & Cells

Parafollicular cells (C cells) – Secrete calcitonin Follicular cells of thyroid – Secrete T3 & T4

- Chief cells of parathyroid Secrete PTH
- Cells most sensitive to hypoxia Neurons
- Fat necrosis Common in breast
- Coagulative necrosis Common in heart, liver, kidney (not brain)

Embryology

- Congenital hydrocele Due to patent processus vaginalis
- Hypertrophic pyloric stenosis Congenital hypertrophy of pyloric muscle

High-Yield Pathology Points

1. Cellular Adaptations & Injury

- Atrophy: Decreased cell size (e.g., disuse atrophy, denervation).
- **Hypertrophy:** Increased cell size (e.g., LVH in hypertension, uterine hypertrophy in pregnancy).
- **Hyperplasia:** Increased cell number (e.g., endometrial hyperplasia, BPH).
- **Metaplasia:** Replacement of one cell type with another (e.g., Barrett's esophagus, squamous metaplasia in smokers).
- **Dysplasia:** Disordered growth (premalignant).
- Necrosis vs. Apoptosis:
 - □ **Necrosis:** Inflammatory, pathological (coagulative, liquefactive, caseous, fat, gangrenous).
 - ☐ **Apoptosis:** Programmed, non-inflammatory (e.g., embryogenesis, viral hepatitis).

2. Inflammation & Repair

Acute Inflammation:

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		☐ Cardinal signs: Redness, swelling, heat, pain, loss of function.
		☐ Mediators: Histamine (immediate), prostaglandins (pain/fever), bradykinin
		(pain).
		Cells: Neutrophils (first responders).
	•	Chronic Inflammation:
		☐ Cells: Macrophages, lymphocytes, plasma cells.
		☐ Granuloma: Caseating (TB) vs. non-caseating (sarcoidosis).
	•	Healing:
		☐ Primary intention: Clean surgical wounds.
		☐ Secondary intention: Large wounds with granulation tissue.
3.	Н	emodynamic Disorders
	•	Edema:
		\Box Causes: \uparrow Hydrostatic pressure (CHF), \downarrow Oncotic pressure (nephrotic syndrome).
	•	Thrombosis: Virchow's triad (stasis, hypercoagulability, endothelial injury).
	•	Embolism:
		□ DVT → PE (most common).
		☐ Fat embolism (long bone fractures).
		☐ Amniotic fluid embolism (DIC, hypoxia during labor).
	•	Infarction:
		☐ Pale (arterial occlusion, e.g., MI).
		☐ Hemorrhagic (venous occlusion, e.g., testicular torsion).
4.	Im	nmunopathology
	•	Hypersensitivity Reactions:
		☐ Type I (IgE): Anaphylaxis, asthma.
		☐ Type II (IgG/IgM): Autoimmune hemolytic anemia, Goodpasture's.
		☐ Type III (Immune complex): SLE, post-streptococcal GN. ☐ Type
		IV (T-cell): TB, contact dermatitis.
	•	Autoimmune Diseases:
		☐ SLE: Anti-dsDNA, anti-Smith.
		☐ Rheumatoid arthritis: RF, anti-CCP.

• Amyloidosis: Congo red (+) with apple-green birefringence.

5. Neoplasia

Benign vs. Malignant:



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	Benign: Well-differentiated, no metastasis.
	☐ Malignant: Poorly differentiated, invades/metastasizes.
	Carcinogenesis:
	☐ Oncogenes: RAS, MYC, HER2/neu.
	☐ Tumor suppressors: p53, Rb, BRCA1/2.
	Paraneoplastic Syndromes:
	☐ SIADH (small cell lung cancer).
	☐ Cushing's (ACTH-secreting tumors).
	☐ Hypercalcemia (PTHrp in squamous cell carcinoma).
6.	Genetic Disorders
	Autosomal Dominant: Huntington's, Marfan's, NF1.
	Autosomal Recessive: Cystic fibrosis, sickle cell, PKU.
	X-linked: Duchenne MD, hemophilia A/B.
	 Triplet Repeat Disorders: Fragile X (CGG), Huntington's (CAG).
7.	Nutritional & Environmental Pathology
	Vitamin Deficiencies:
	☐ B1 (Thiamine): Beriberi, Wernicke-Korsakoff.
	☐ B3 (Niacin): Pellagra (3 D's: Dermatitis, Diarrhea, Dementia).
	☐ B12/Folate: Megaloblastic anemia, neurologic symptoms (B12 only).
	☐ Vitamin C: Scurvy (gingival bleeding, poor wound healing).
	Toxins:
	☐ CO poisoning: Cherry-red skin, ↑ COHb.
	☐ Lead poisoning: Basophilic stippling, wrist/foot drop.
8.	Infectious Disease Pathology
	• Bacteria:
	☐ Staph aureus: Abscesses, endocarditis (IV drug users).
	☐ Strep pyogenes: Rheumatic fever (M protein), post-streptococcal GN.
	• Viruses:
	☐ HPV : Koilocytes, cervical cancer (types 16/18).

☐ EBV: Mononucleosis (atypical lymphocytes), Burkitt's lymphoma. Fungi: Candida: Pseudohyphae, thrush (immunocompromised). ☐ Aspergillus: Angioinvasive (neutropenic patients).
9. Hematopathology
Anemias:
\square Microcytic: Fe deficiency (\downarrow ferritin), thalassemia (\uparrow HbA2).
☐ Macrocytic: B12/folate deficiency (hypersegmented neutrophils). ☐
Hemolytic: G6PD deficiency (Heinz bodies, bite cells).
• Leukemia/Lymphoma:
☐ AML: Auer rods, myeloperoxidase (+).
☐ CLL: Smudge cells, CD5/CD19 (+).
☐ Hodgkin lymphoma: Reed-Sternberg cells (CD15/CD30 +).
10. Systemic Pathology Highlights
• Cardiac:
\square Atherosclerosis: LDL \rightarrow foam cells \rightarrow fibrous plaque \rightarrow rupture.
☐ MI complications: Papillary muscle rupture (MR), ventricular aneurysm.
Pulmonary:
\square Emphysema: $\alpha 1$ -antitrypsin deficiency (panacinar).
☐ ARDS: Diffuse alveolar damage, hyaline membranes.
• Renal:
☐ Nephritic syndrome: HTN, hematuria (post-streptococcal GN).
☐ Nephrotic syndrome: Proteinuria, hypoalbuminemia (minimal change disease)
• Liver:
☐ Cirrhosis: Bridging fibrosis, portal HTN.
☐ Hepatitis:
Acute: Councilman bodies (apoptosis).
☐ Chronic: Ground-glass hepatocytes (HBV).

Key Mnemonics

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• Granulomatous diseases: TB, Sarcoidosis, Crohn's, Fungal, Beryllium (TSCFB).

Tumor markers:

CA-125 (ovary), PSA (prostate), CEA (colon).

 Hypersensitivity reactions: ACID (Type I – Anaphylactic, Type II – Cytotoxic, Type III – Immune complex, Type IV – Delayed).

High-Yield Notes from Anatomy Shelf Notes

1. Lumbar Puncture & Epidural Anesthesia

- Entry Point: Between L3/L4 or L4/L5 (level of iliac crests).
- **Structures Pierced:** Skin → Fascia → Ligamentum flavum → Epidural space → Dura mater → Subarachnoid space (CSF).
- Spinal Cord Ends:

Adults: L1/L2.

o Children: L3.

- Dural Sac Ends: S2.
- **Complication:** Herniation if intracranial pressure is elevated.

2. Herniated IV Disc

- Common Sites: L4/L5 or L5/S1 (lumbar); C5/C6 or C6/C7 (cervical).
- Affected Nerve Root: Traversing root (e.g., L4/L5 herniation compresses L5).
- **Mechanism:** Nucleus pulposus herniates through anulus fibrosus, compressing spinal nerve.

3. Spinal Curvature Abnormalities

- Kyphosis: Exaggerated thoracic curve (osteoporosis in elderly).
- Lordosis: Exaggerated lumbar curve (pregnancy, spondylolisthesis).
- Scoliosis: Lateral deviation (leg-length discrepancy, poliomyelitis).

4. Upper Limb Fractures

Humerus Fractures:

o Surgical Neck: Axillary nerve injury. o Midshaft: Radial nerve injury.

- o Medial Epicondyle: Ulnar nerve injury.
- Colles' Fracture: Distal radius fracture with dorsal displacement ("dinner fork deformity").
- Scaphoid Fracture:
 - Tenderness in anatomical snuffbox.
 - Risk of avascular necrosis (proximal fragment).

5. Rotator Cuff Muscles (SITS)

- Supraspinatus, Infraspinatus, Teres minor, Subscapularis.
- Function: Stabilize shoulder joint; supraspinatus initiates abduction (0°-15°).

6. Carpal Tunnel Syndrome

- Cause: Compression of median nerve in carpal tunnel.
- Symptoms: O Numbness in lateral 3.5 fingers (spares palm).
 - Ape hand deformity (loss of opposition).

7. Brachial Plexus Injuries

- **Erb-Duchenne Palsy (C5–C6):** o "Waiter's tip" posture (adducted shoulder, medially rotated arm).
- Klumpke Palsy (C8–T1):
 - o Claw hand (ulnar nerve) + ape hand (median nerve).

8. Lower Limb Injuries

- Femoral Neck Fracture: Risk of avascular necrosis (disrupted retinacular arteries).
- Unhappy Triad (Knee):
 - o Tibial collateral ligament + Medial meniscus + Anterior cruciate ligament (ACL).
- Ankle Sprain: Inversion injury damages anterior talofibular ligament.

9. Abdominal Hernias

- Indirect Inguinal Hernia: O Congenital; passes lateral to inferior epigastric vessels.
- Direct Inguinal Hernia:
 - o Acquired; passes **medial** to inferior epigastric vessels (Hesselbach's triangle).

10. Gastrointestinal Anatomy

- Foregut/Midgut/Hindgut:
 - **Foregut:** Supplied by **celiac artery** (T5–T9); pain referred to **epigastrium**. **Midgut:** Supplied by **SMA** (T10–T11); pain referred to **umbilicus**.
 - Hindgut: Supplied by IMA (L1–L2); pain referred to hypogastrium.
- Appendicitis:
 - o Early pain at umbilicus (visceral); later shifts to McBurney's point (parietal).

11. Portal Hypertension

- Portacaval Anastomoses:
 - Esophageal varices (left gastric vein + azygos vein).
 - o Caput medusae (paraumbilical veins + epigastric veins).
 - Hemorrhoids (superior rectal vein + inferior rectal vein).

12. Cranial Nerves

- Facial Nerve (CN VII):
 - Bell's Palsy: Ipsilateral facial paralysis, dry eye, loss of taste (anterior 2/3 tongue).
- Oculomotor Nerve (CN III):
 - o **Palsy:** Ptosis, "down and out" eye, dilated pupil.
- Horner Syndrome:
 - Miosis, ptosis, anhydrosis (disrupted sympathetic pathway).

13. Thyroid & Parathyroid

- Thyroid:
 - Superior laryngeal nerve (external branch) at risk during thyroidectomy.

- o **Recurrent laryngeal nerve** injury → hoarseness.
- Parathyroid:
 - o **PTH** increases blood calcium; **calcitonin** decreases it.

14. Larynx

- Vocal Cord Innervation:
 - o **Recurrent laryngeal nerve** (all muscles except cricothyroid).
 - o **Cricothyroid muscle** (external laryngeal nerve).

15. Emergency Procedures

• Cricothyrotomy: Incision through cricothyroid membrane for airway obstruction.

Key Mnemonics

- Rotator Cuff: SITS (Supraspinatus, Infraspinatus, Teres minor, Subscapularis).
- Foregut/Midgut/Hindgut: CA (celiac), SMA, IMA.
- Brachial Plexus Roots: "Randy Travis Drinks Cold Beer" (Roots, Trunks, Divisions, Cords, Branches).

High-Yield Revision Points Gynae and Obs

1. Pregnancy & Labour

•	rerminology:
	☐ Term pregnancy: 37–42 weeks
	☐ Preterm labour: <37 weeks ☐ Post -
	term pregnancy: >42 weeks
•	Stages of labour:
	☐ 1st stage: Cervical dilation (latent & active phases)
	☐ 2nd stage: Full dilation to delivery

	☐ 3rd stage: Placental delivery (expectant vs. active management)
•	Induction of labour:
	☐ Indications: Post-term, preeclampsia, IUGR
	☐ Methods: Prostaglandins (PGE2), oxytocin, ARM
М	ledical Disorders in Pregnancy
•	Hypertensive disorders:
	☐ Gestational HTN: New HTN after 20 weeks, no proteinuria
	☐ Preeclampsia: HTN + proteinuria/organ dysfunction
	☐ Eclampsia: Preeclampsia + seizures (treat with MgSO₄) ☐
	HELLP syndrome: Hemolysis, elevated LFTs, low platelets
•	Diabetes in pregnancy:
	☐ GDM: Screen at 24–28 weeks (OGTT)
	□ Management: Diet → insulin (oral agents avoided)
•	Thromboembolism:
	☐ LMWH is anticoagulant of choice
Λ./	alpresentation & Multiple Pregnancy
IVI	ulpresentation & whitiple Fregnancy
•	Breech presentation:
	☐ Types: Frank, complete, footling
	☐ Management: ECV at 37 weeks → C-section if unsuccessful
•	Transverse lie:
	☐ Risk factors: Multiparity, polyhydramnios
	☐ Delivery: C-section (vaginal delivery contraindicated)
•	Twin pregnancy:
	☐ Dichorionic diamniotic (DCDA): Lowest risk
	☐ Monochorionic monoamniotic (MCMA): Highest risk (cord entanglement)
	☐ Delivery: Vaginal if Twin A cephalic, C-section if complications
Lic	quor Volume Abnormalities
•	Polyhydramnios (AFI >24 cm):
	☐ Causes: GDM, fetal anomalies (e.g., duodenal atresia)
•	Oligohydramnios (AFI <5 cm):

2.

3.

4.

	□ Causes: PPROM, IUGR, renal anomalies□ Management: Amnioinfusion if severe
5.	Obstetric Emergencies
	Placental abruption:
	☐ Classic triad: Painful bleeding, uterine tenderness, fetal distress
	☐ Management: Emergency delivery
	• Uterine rupture:
	☐ Risk factors: Previous C-section, uterine surgery ☐ Signs:
	Sudden pain, fetal distress, loss of contractions
	Shoulder dystocia:
	☐ McRoberts maneuver (first-line)
	☐ Complication: Erb's palsy
G۱	necology Topics
-,	Andrew Spires
<i>6</i> .	Puberty
	Normal puberty:
	☐ Thelarche (breast buds) → Pubarche → Menarche
	\square Delayed puberty: No breast development by 13 or no menarche by
	Primary amenorrhea:
	☐ Turner syndrome (45X): Streak ovaries, high FSH
	☐ Imperforate hymen: Cyclical pain, bulging membrane
7.	Infertility
	• Causes:
	☐ Male factor: Low sperm count/motility
	☐ Female factor: PCOS, tubal blockage, endometriosis
	• Investigations:
	☐ Day 3 FSH/LH, HSG, semen analysis
	1 buy 5 1 511, 1150, semen unarysis
	• Treatment:

8. Contraception

	•	Combined hormonal (COCP):
		☐ Contraindications: HTN, smokers >35, history of VTE
	•	Progesterone-only (POP):
		☐ Safe in breastfeeding, VTE risk
	•	LARC (Long-acting reversible):
		☐ Copper IUD: Emergency contraception
		☐ Levonorgestrel IUS (Mirena): Treats heavy bleeding
9.	M	enstrual Disorders
	•	Heavy menstrual bleeding (HMB):
		☐ Causes: Fibroids, adenomyosis, coagulopathy (e.g., von Willebrand)
		☐ Treatment: Tranexamic acid, hormonal IUD, endometrial ablation
	•	Dysmenorrhea:
		☐ Primary: No pathology (treat with NSAIDs)
		☐ Secondary: Endometriosis, PID
10.	. Se.	xual Disorders
	•	Dyspareunia (painful sex):
		☐ Superficial: Vaginismus, vulvodynia
		☐ Deep: Endometriosis, PID •
	Ну	poactive sexual desire disorder:
		☐ Postpartum: Low estrogen (breastfeeding)
11.	. Ur	inary Problems
	•	Stress incontinence:
		□ Pelvic floor exercises (1st line) → Midurethral sling
	•	Overactive bladder:
		☐ Anticholinergics (oxybutynin), mirabegron

12. Uterovaginal Prolapse

•	Grades:
	☐ 1st: Descent to introitus
	☐ 2nd: Protrudes outside ☐
	3rd/4th: Complete procidentia
•	Management:
	□ Pessary (1st line in elderly) → Surgery (hysterectomy + repair)
13. Ge	nital Tract Infections
•	Bacterial vaginosis:
	☐ Clue cells, fishy odor → Metronidazole
•	Candidiasis:
	☐ Itching, cottage cheese discharge → Fluconazole
•	PID:
	□ Cervical motion tenderness → Ceftriaxone + Doxycycline
14. Gy	necological Tumors
•	Endometrial cancer:
	□ Postmenopausal bleeding → Hysterectomy
•	Ovarian cancer:
	☐ Silent tumor → CA-125, ultrasound
	□ Risk factors: BRCA Lynch syndrome

Key Mnemonics

- HELLP syndrome: Hemolysis, Elevated LFTs, Low Platelets
- **PID treatment: CDC (C**eftriaxone, **D**oxycycline, **C**over anaerobes with metronidazole if abscess)
- Polyhydramnios causes: Diabetes, Duodenal atresia, Down syndrome

Internal Medicine

Endocrinology

1. Pituitary Gland

- Anterior Pituitary (Adenohypophysis):
 - Prolactinoma: Most common functional pituitary adenoma → galactorrhea, amenorrhea, infertility (treated with dopamine agonists like bromocriptine/cabergoline).
 - Growth Hormone (GH) Excess:
 - → Before epiphyseal closure: Gigantism.
 - ★ After closure: Acromegaly (enlarged hands/feet, coarse facial features, macroglossia, carpal tunnel syndrome).
 - → Diagnosis: Elevated IGF-1, failure to suppress GH on oral glucose tolerance test (OGTT). ○ Cushing's Disease: ACTH-secreting adenoma → high cortisol (diagnose with dexamethasone suppression test). ○ Sheehan's Syndrome: Postpartum pituitary necrosis (hypotension + failure to lactate + amenorrhea).
- Posterior Pituitary (Neurohypophysis):
 - Diabetes Insipidus (DI):
 - → Central DI: ADH deficiency (head trauma, tumors) → low urine osmolality, high serum osmolality (treat with desmopressin (dDAVP)).
 - **→** Nephrogenic DI: ADH resistance (lithium, hypercalcemia) → no response to dDAVP (treat with thiazides + low-salt diet). SIADH: High ADH → hyponatremia, high urine osmolality, low serum osmolality (treat with fluid restriction, demeclocycline).

2. Thyroid Disorders

- Hyperthyroidism:
 - ⊙ Graves' Disease: TSI antibodies → diffuse goiter, exophthalmos, pretibial myxedema (treat with methimazole/propylthiouracil (PTU) or radioactive iodine).
 - Thyroid Storm: Tachycardia, fever, agitation (treat with PTU (blocks synthesis + conversion), beta-blockers, steroids, iodine).
 - Toxic Multinodular Goiter: Hot nodules on scan (no autoantibodies).
- Hypothyroidism:

- Hashimoto's Thyroiditis: Anti-TPO antibodies → goiter, weight gain,
 bradycardia, myxedema coma (treat with levothyroxine). Myxedema Coma:
 Hypothermia, bradycardia, coma (treat with IV levothyroxine + steroids).
- Thyroid Nodules/Cancer:
 - Most common type: Papillary carcinoma (Psammoma bodies, lymphatic spread).
 Medullary carcinoma: Calcitonin secretion, associated with MEN 2A/2B (RET proto-oncogene).
 - Anaplastic carcinoma: Aggressive, poor prognosis.

3. Adrenal Gland

- Cushing's Syndrome:
 - Causes: Pituitary adenoma (Cushing's disease), adrenal adenoma, ectopic ACTH (small cell lung cancer), exogenous steroids. O Findings: Moon facies, buffalo hump, striae, hyperglycemia, osteoporosis.
 - Diagnosis: High-dose dexamethasone suppression test (suppresses if pituitary source).
- Primary Hyperaldosteronism (Conn's Syndrome):
 - o **Hypokalemia, metabolic alkalosis, hypertension** (low renin, high aldosterone).
 - Diagnosis: Elevated aldosterone/renin ratio (treat with spironolactone).
- Addison's Disease (Primary Adrenal Insufficiency):
 - Causes: Autoimmune (most common), TB, metastatic disease.
 - Findings: Fatigue, hyperpigmentation, hyponatremia, hyporkalemia, hypoglycemia.
 - Diagnosis: Low cortisol, high ACTH (treat with glucocorticoids + mineralocorticoids).
- Pheochromocytoma: O Rule of 10s: 10% bilateral, 10% malignant, 10% extra-adrenal. O Findings: Episodic hypertension, headache, sweating, palpitations.
 - O Diagnosis: 24-hour urine metanephrines (treat with alpha-blockers first (phenoxybenzamine), then beta-blockers).

4. Diabetes Mellitus (DM)

- Type 1 DM: Autoimmune (anti-GAD antibodies), absolute insulin deficiency, DKA risk.
- Type 2 DM: Insulin resistance, associated with metabolic syndrome (obesity, HTN, dyslipidemia).

Complications:

- Microvascular: Retinopathy, nephropathy (Kimmelstiel-Wilkin nodules), neuropathy.
- Macrovascular: CAD, stroke, PVD. DKA: High anion gap metabolic acidosis, ketones, Kussmaul breathing (treat with IV fluids, insulin, potassium).
- HHS: Hyperosmolar state without significant ketosis (higher mortality than DKA).

5. Calcium & Bone Disorders

- Hyperparathyroidism:
 - Primary: High PTH, high calcium (adenoma) → osteitis fibrosa cystica, nephrolithiasis.
 - o Secondary: High PTH, low calcium (chronic kidney disease).
- Hypoparathyroidism: Low PTH, low calcium (post-thyroidectomy) → Chvostek's & Trousseau's signs.
- **Hypercalcemia of Malignancy**: **PTHrP secretion** (squamous cell lung cancer, breast cancer).
- Osteoporosis: DEXA scan (T-score ≤ -2.5), treat with bisphosphonates.

6. Reproductive Endocrinology

- PCOS: Oligomenorrhea, hirsutism, insulin resistance, LH:FSH > 2:1.
- Klinefelter's (47,XXY): Tall, gynecomastia, small testes, infertility.
- Turner's (45,X): Short stature, webbed neck, coarctation of aorta, primary amenorrhea.

7. Miscellaneous

- MEN Syndromes:
 O MEN 1 (3 P's): Pituitary, Pancreatic (gastrinoma), Parathyroid.
 - MEN 2A: Medullary thyroid cancer, Pheochromocytoma, Hyperparathyroidism.
 MEN 2B: Medullary thyroid cancer, Pheochromocytoma, Marfanoid habitus, mucosal neuromas.

Key Labs & Diagnostic Tests

- ACTH Stimulation Test: Diagnoses adrenal insufficiency.
- Water Deprivation Test: Differentiates DI vs. psychogenic polydipsia.
- Dexamethasone Suppression Test: Diagnoses Cushing's.

Gastroenterology

1. Esophagus

GERD:

- o **Symptoms**: Heartburn, regurgitation, chronic cough, hoarseness.
- Complications: Barrett's esophagus (intestinal metaplasia → adenocarcinoma), strictures.
- Diagnosis: Endoscopy (gold standard), pH monitoring. Treatment: PPIs (1st-line), H2 blockers, lifestyle changes (avoid fatty foods, caffeine, alcohol).

Achalasia:

- o Pathology: Loss of myenteric (Auerbach's) plexus \rightarrow failure of LES relaxation.
- Symptoms: Dysphagia (solids > liquids), regurgitation of undigested food, bird's beak on barium swallow.
- Diagnosis: Esophageal manometry (gold standard).
- o Treatment: Pneumatic dilation, Heller myotomy, Botox injection.

Boerhaave Syndrome:

- Full-thickness esophageal rupture (after vomiting) → mediastinitis, subcutaneous emphysema, Hamman's crunch.
- Treatment: Emergency surgery + antibiotics.

2. Stomach & Peptic Ulcer Disease (PUD)

PUD Causes:

○ **H. pylori** (most common, **urea breath test** for diagnosis). ○ **NSAIDs** (COX-1 inhibition $\rightarrow \downarrow$ PGE2 $\rightarrow \downarrow$ mucosal protection). ○ **Zollinger-Ellison Syndrome** (gastrinoma \rightarrow high acid secretion \rightarrow multiple ulcers).

Complications:

- Bleeding (most common).
- Perforation: Sudden severe abdominal pain, rigid abdomen, free air under diaphragm on X-ray.
- o Gastric Outlet Obstruction: Non-bilious vomiting, succussion splash.

Treatment:

- H. pylori: PPI + clarithromycin + amoxicillin/metronidazole (triple therapy).
- NSAID-induced: Stop NSAIDs + PPI.

3. Liver Disease

- Cirrhosis & Portal Hypertension:
 - Causes: Alcohol, hepatitis B/C, NASH.
 - Complications:
 - **+ Esophageal varices** (treat acute bleed with **octreotide + band ligation**).
 - Ascites (diagnose with SAAG >1.1, treat with spironolactone + furosemide).
 - → Hepatic encephalopathy (↑ammonia, treat with lactulose + rifaximin).
 - → Hepatorenal syndrome (oliguria, ↑Cr, treat with midodrine + octreotide).

Hepatitis:

- Hep B: HBsAg (active infection), Anti-HBc (exposure), Anti-HBs (immunity).
- Hep C: Most common cause of chronic hepatitis → cirrhosis/HCC (treat with direct-acting antivirals).

Liver Tumors:

- Hepatocellular Carcinoma (HCC): ↑AFP, associated with cirrhosis, hepatitis B/C.
- Metastases: Most common liver tumor (colon, breast, lung primaries).

4. Biliary & Pancreatic Disease

• Gallstones:

- Cholelithiasis: Asymptomatic. O Cholecystitis: RUQ pain, Murphy's sign, fever (treat with cholecystectomy).
- Choledocholithiasis: Jaundice, ↑ALP, ↑bilirubin (diagnose with MRCP, treat with ERCP).

Acute Pancreatitis:

- Causes: GET SMASHED (Gallstones, Ethanol, Trauma, Steroids, Mumps, Autoimmune, Scorpion sting, Hyperlipidemia, ERCP, Drugs). ○ Diagnosis:
 ↑Lipase (more specific than amylase), CT findings.
- Complications: Pseudocyst (4 weeks later), necrotizing pancreatitis.
- Chronic Pancreatitis:

- Causes: Alcohol most common.
- Findings: Steatorrhea (fat malabsorption), calcifications on X-ray, DM (loss of islets).

5. Inflammatory Bowel Disease (IBD)

- Crohn's Disease:
 - Transmural inflammation, skip lesions, cobblestone mucosa, fistulas, noncaseating granulomas.

 Extraintestinal: Erythema nodosum, uveitis, ankylosing spondylitis (HLAB27).
 - Treatment: Steroids, anti-TNF (infliximab), surgery (no cure).
- Ulcerative Colitis:
 - o Limited to colon, continuous inflammation, crypt abscesses, no granulomas.
 - o Extraintestinal: Primary sclerosing cholangitis (PSC), pyoderma gangrenosum.
 - Toxic Megacolon: Medical emergency (abdominal distension, fever, tachycardia).
 - Treatment: 5-ASA (mesalamine), colectomy (curative).

6. Colon & Small Bowel Disorders

- Diverticulosis:
 - Asymptomatic outpouchings, risk ↑ with age.
 - Diverticulitis: LLQ pain, fever, leukocytosis (treat with antibiotics).
- Celiac Disease:
 - Autoimmune (anti-tTG, anti-endomysial antibodies), villous atrophy, diarrhea, bloating.
 - o Treatment: Gluten-free diet.
- Colorectal Cancer:
 - Most common site: Rectosigmoid.
 - Screening: Colonoscopy at 45+ (earlier if familial polyposis, Lynch syndrome).

7. GI Bleeding

- Upper GI Bleed (UGIB):
 - Causes: PUD, varices, Mallory-Weiss tear. Diagnosis: EGD.
- Lower GI Bleed (LGIB):

- Causes: Diverticulosis (most common), angiodysplasia, hemorrhoids.
- Diagnosis: Colonoscopy.

8. Miscellaneous High-Yield Facts

- Appendicitis: RLQ pain, McBurney's tenderness, ↑WBC, Rovsing's sign.
- Hereditary Hemochromatosis: ↑Fe, ↑ferritin, ↑TSAT, cirrhosis, DM, cardiomyopathy.
- Wilson's Disease: ↓Ceruloplasmin, Kayser-Fleischer rings, neuropsych symptoms.

Key Labs & Imaging

- AST/ALT Ratio: >2 in alcoholic hepatitis.
- Ammonia: ↑ in hepatic encephalopathy.
- Barium Swallow: Bird's beak (achalasia), corkscrew (esophageal spasm).

Neurology

1. Stroke & Cerebrovascular Disease

- Ischemic Stroke (80%):
 - Thrombotic: Atherosclerosis (MCA most common). Embolic: Atrial fibrillation (cardioembolic), MCA territory most affected.
 - Lacunar Strokes: Small vessel disease (hypertension, diabetes) → pure motor (internal capsule), pure sensory (thalamus), ataxic hemiparesis (pons). ○
 Diagnosis: Non-contrast CT first (rule out hemorrhage), MRI diffusionweighted imaging (DWI) gold standard. ○ Treatment: tPA within 4.5 hrs (exclude hemorrhage, recent surgery, anticoagulation).
- Hemorrhagic Stroke (20%):

Hypertensive hemorrhage: Basal ganglia, thalamus, pons, cerebellum.

- Subarachnoid Hemorrhage (SAH): Thunderclap headache, nuchal rigidity,
 Fisher Grade 3 = highest vasospasm risk.
 - ★ Cause: Berry aneurysm (most commonly at anterior communicating artery).
 - **→** Diagnosis: CT **→** LP if negative (xanthochromia).
 - **→** Treatment: Nimodipine (prevents vasospasm), surgical clipping/coiling.
- Transient Ischemic Attack (TIA):
 - Neurologic deficits resolving <24 hrs (usually <1 hr).
 - ABCD2 score predicts stroke risk.

2. Seizures & Epilepsy

- Focal Seizures:
 - With impaired awareness (complex partial): Temporal lobe (olfactory/psychic auras, automatisms).
 Without impaired awareness: Motor (Jacksonian march) or sensory symptoms.
- Generalized Seizures:
 - Tonic-clonic (grand mal): Postictal confusion, tongue biting. O Absence (petit mal): 3 Hz spike-and-wave EEG, no postictal state. O Atonic: Sudden loss of tone ("drop attacks").
 - Myoclonic: Sudden jerks (e.g., Juvenile Myoclonic Epilepsy).
- Status Epilepticus: >5 mins of seizure or recurrent without recovery → IV lorazepam, then fosphenytoin.
- First-line drugs:
 - Focal: Levetiracetam, carbamazepine.
 - o **Generalized**: Valproate (avoid in pregnancy \rightarrow neural tube defects).

3. Headache

- Migraine:
 - With aura (scintillating scotoma, paresthesias), photophobia, phonophobia.
 Treatment: Triptans (contraindicated in CAD, stroke), propranolol/topiramate for prophylaxis.
- Cluster Headache:

- Unilateral, periorbital, Horner's syndrome (ptosis, miosis).
- Treatment: High-flow O2, sumatriptan.
- Trigeminal Neuralgia:
 - Unilateral stabbing pain (V2/V3 distribution) \rightarrow carbamazepine.
- Idiopathic Intracranial Hypertension (Pseudotumor cerebri):
 - Obese women, papilledema, headache, visual loss.
 - Treatment: Acetazolamide, LP, weight loss.

4. Neurodegenerative Disorders

- Alzheimer's Disease:
 - Memory loss (early short-term), apraxia, aphasia. O Pathology: Amyloid plaques (Aβ), neurofibrillary tangles (tau).
 - Treatment: AChE inhibitors (donepezil), memantine (NMDA antagonist).
- Parkinson's Disease:
 - TRAP: Tremor (pill-rolling), Rigidity (cogwheel), Akinesia/bradykinesia, Postural instability.
 - Pathology: Lewy bodies (α-synuclein), substantia nigra degeneration.
 - Treatment: Levodopa/carbidopa (gold standard), dopamine agonists (pramipexole).
- ALS (Lou Gehrig's Disease):
 - UMN + LMN signs (spasticity + atrophy, no sensory loss).
 - No cure (riluzole modestly prolongs survival).

5. Demyelinating Diseases

- Multiple Sclerosis (MS):
 - Relapsing-remitting (most common), optic neuritis, internuclear ophthalmoplegia (INO), Lhermitte's sign.
 - Diagnosis: MRI (periventricular plaques), CSF oligoclonal bands.
 - Treatment: Steroids (acute), interferon-β/glatiramer (prevention).
- Guillain-Barré Syndrome:

Ascending paralysis, areflexia, albuminocytologic dissociation (个CSF protein, normal WBC).

Treatment: IVIG or plasmapheresis.

6. CNS Infections

- Meningitis:
 - Bacterial (S. pneumoniae, N. meningitidis): Neck stiffness, fever, Kernig's/Brudzinski's signs → empiric ceftriaxone + vancomycin + dexamethasone.
 - Viral (enterovirus): Lymphocytic CSF, self-limiting.
- Encephalitis: O HSV-1: Temporal lobe necrosis (MRI), acyclovir treatment.
- Brain Abscess:
 - Ring-enhancing lesion, Streptococci/Staphylococci → surgical drainage + antibiotics.

7. Spinal Cord & Peripheral Nerve

- Spinal Cord Compression:
 - o Emergency: Back pain, weakness, sensory level, bowel/bladder dysfunction.
 - o Causes: Metastasis (most common), epidural abscess.
- Brown-Séquard Syndrome:
 - Hemisection: Ipsilateral weakness + loss of vibration/proprioception, contralateral loss of pain/temp.
- Peripheral Neuropathy:
 - Diabetes: Glove-and-stocking distribution. O B12 Deficiency: Subacute combined degeneration (posterior + lateral columns).

8. High-Yield Syndromes

• Horner's Syndrome: Ptosis, miosis, anhidrosis (Pancoast tumor, carotid dissection).

- Wernicke-Korsakoff: Confusion, ataxia, nystagmus (Wernicke) + confabulation (Korsakoff) → thiamine deficiency.
- Syringomyelia: Bilateral loss of pain/temp in "cape-like" distribution (central cord lesion).

Key Reflexes & Localization

- Babinski Sign: UMN lesion (toes dorsiflex).
- LMN Lesion: Hyporeflexia, atrophy, fasciculations.
- UMN Lesion: Hyperreflexia, spasticity, clasp-knife rigidity.

Must-Know Drugs & Toxins

- Warfarin Toxicity: Vitamin K + FFP.
- Opioid Overdose: Pinpoint pupils, naloxone reversal.
- Anticholinergic Toxicity: Hot as a hare, dry as a bone, red as a beet, mad as a hatter.

Cardiology

1. Coronary Artery Disease (CAD) & Acute Coronary Syndrome (ACS)

- Stable Angina:
 - Symptoms: Chest pain (crushing, substernal) radiating to left arm/jaw, relieved by rest/nitroglycerin.
 - Diagnosis: Stress test (exercise ECG), coronary angiography (gold standard).
 Treatment: Nitrates, beta-blockers (1st-line), CCBs (if beta-blockers contraindicated).
- Unstable Angina/NSTEMI:
 - **Symptoms**: Angina at rest, **↑Troponin (NSTEMI)**, normal **ST on ECG**.
 - Treatment: Aspirin + P2Y12 inhibitor (clopidogrel), heparin, statin, early invasive strategy (PCI).

STEMI:

- ECG Findings: ST elevation (>1mm in 2+ contiguous leads), reciprocal ST depression.
- Treatment: Reperfusion ASAP → PCI (preferred) or fibrinolytics (tPA if PCI >90 min away).
- o Complications:
 - → Ventricular arrhythmias (VF/VT) → defibrillation.
 - **+** Cardiogenic shock (↓BP, pulmonary edema) **→** inotropes (dobutamine), IABP.
 - → Papillary muscle rupture → acute mitral regurgitation (holosystolic murmur).

2. Heart Failure (HF)

- Systolic HF (HFrEF, EF <40%):
 - Causes: Ischemic cardiomyopathy, dilated cardiomyopathy (DCM).
 - Treatment: ACEI/ARB + beta-blocker (carvedilol/metoprolol) + spironolactone
 + SGLT2 inhibitors.
- Diastolic HF (HFpEF, EF ≥50%):
 - Causes: HTN, hypertrophic cardiomyopathy (HCM), amyloidosis.
 - Treatment: Diuretics (symptom relief), BP control.
- Acute Decompensated HF:
 - O Symptoms: Dyspnea, crackles, S3 gallop, JVD, peripheral edema.
 - Treatment: Diuretics (furosemide), vasodilators
 (nitroprusside/nitroglycerin), inotropes (dobutamine if hypotensive).

3. Valvular Heart Disease

V		
al	Stenosis	Regurgitation
ve		

A or ti c	Systolic ejection murmur (SEM), syncope, angina, S4	Early diastolic decrescendo murmur, wide pulse pressure, bounding pulses
M itr al	Diastolic rumble, opening snap, LA enlargement (afib)	Holosystolic murmur at apex, radiating to axilla

- Aortic Stenosis: Triad → Syncope, angina, dyspnea.
- Mitral Regurgitation: Acute (papillary muscle rupture) → pulmonary edema.
- Mitral Valve Prolapse: Mid-systolic click + late systolic murmur, anxiety, tall/thin body habitus.

4. Arrhythmias

- Atrial Fibrillation (A-fib):
 - o Irregularly irregular pulse, no P waves.
 - o Treatment:
 - **→ Rate control** (beta-blockers, CCBs, digoxin).
 - Rhythm control (amiodarone, cardioversion if unstable). →
 Anticoagulation (CHADS₂-VASc ≥2 → warfarin/DOACs).
- Ventricular Tachycardia (VT):
 - Wide QRS, no P waves. Treatment: Unstable → cardioversion, stable → amiodarone/lidocaine.

Bradycardia:

- Sinus node dysfunction (sick sinus syndrome) → pacemaker.
- O AV Blocks:
 - **→** 1st-degree: PR >200ms.
 - **→** 2nd-degree (Type I: Wenckebach): PR prolongation **→** dropped QRS.
 - **+** 3rd-degree (Complete): P waves and QRS dissociated → pacemaker.

5. Hypertension (HTN)

- Essential (Primary) HTN (90%): No identifiable cause.
- Secondary HTN Causes:
 - Renal artery stenosis → ↑Renin, unilateral small kidney.
 Pheochromocytoma → Episodic HTN, ↑catecholamines.
 - Cushing's \rightarrow \uparrow Cortisol, moon facies.
- Hypertensive Emergency: BP >180/120 + end-organ damage (encephalopathy, renal failure, papilledema) → IV nitroprusside/labetalol.

6. Cardiomyopathies

Туре	Key Features	Treatment
Dilated (DCM)	↓EF, S3, global hypokinesis	ACEI, beta-blockers
Hypertrophic (HCM)	LV hypertrophy, diastolic dysfunction, systolic murmur 个 with Valsalva	Beta-blockers, avoid nitrates/diuretics
Restrictive (Amyloidosis)	个LV wall thickness, "sparkling" myocardium on echo	Supportive, poor prognosis

7. Pericardial Disease

- Acute Pericarditis:
 - Chest pain improved by sitting forward, pericardial friction rub.
 - ECG: Diffuse ST elevation, PR depression.
 - Treatment: NSAIDs + colchicine.

Cardiac Tamponade:

- Beck's Triad: Hypotension, JVD, muffled heart sounds. O Pulsus paradoxus (>10 mmHg drop in BP with inspiration).
- Treatment: Pericardiocentesis (emergency).

8. Congenital Heart Disease

Defect	Shunt Direction	Key Features
ASD	L → R	Fixed split S2, pulmonary flow murmur
VSD	L → R	Holosystolic murmur at LSB
PDA	L → R	Continuous "machine-like" murmur
Tetralogy of Fallot	R → L	4 features: VSD, RVH, overriding aorta, PS → "Cyanotic spells"

9. High-Yield Pharmacology

- Beta-Blockers: ↓HR, ↓BP (avoid in asthma, decompensated HF).
- CCBs:
 - Dihydropyridines (amlodipine) → vasodilation.
 - Non-DHP (verapamil/diltiazem) → ↓HR, avoid in HF.
- Digoxin: ↑Contractility, ↓HR (toxicity → nausea, yellow vision, arrhythmias).
- Nitrates: Venodilation → ↓preload (contraindicated in RV infarction, HCM).

10. Key ECG Findings

- STEMI: ST elevation.
- Hyperkalemia: Peaked T waves → widened QRS → sine wave → VF.
- Hypokalemia: U waves, flat T waves.
 - Long QT Syndrome: Risk of Torsades de pointes (treat with Mg²⁺).

Nephrology

1. Acid-Base Disorders

Metabolic Acidosis (pH <7.35, HCO₃ - <22)

- Anion Gap (AG) = Na⁺ (Cl₊ + HCO₃) ↑AG (MUDPILES):
 - → Methanol, Uremia, DKA, Paraldehyde, INH/Iron, Lactic

acidosis, Ethylene glycol, Salicylates. O Normal AG (Hyperchloremic):

- **→** Diarrhea, RTA (Type 1-4), acetazolamide, saline infusion.
- Treatment: Correct underlying cause (e.g., insulin for DKA, bicarb for severe acidosis).

Metabolic Alkalosis (pH >7.45, HCO₃ _ >26)

- Causes:
 - Vomiting/NG suction (loss of H^+) \rightarrow hypochloremic, hypokalemic. Hyperaldosteronism ($\uparrow H^+$ excretion).
- Treatment: NS for volume depletion, KCl for hypokalemia, acetazolamide if volume overloaded.

Respiratory Acidosis/Alkalosis

- Respiratory Acidosis (↑pCO₂): COPD, opioid overdose.
- Respiratory Alkalosis (↓pCO₂): Anxiety, PE, salicylate toxicity.

2. Electrolyte Disorders

Hyperkalemia (K+ >5.0)

- Causes: K+ supplements, ACEIs/ARBs, spironolactone, rhabdo, acidosis.
- ECG: Peaked T waves → widened QRS → sine wave → VF.
- Treatment:

Stabilize myocardium: Calcium gluconate. ○ Shift K+ into cells: Insulin + glucose, albuterol, bicarb. ○ Remove K+: Kayexalate, dialysis.

Hypokalemia ($K^+ < 3.5$)

- Causes: Diuretics, vomiting, hyperaldosteronism.
- ECG: U waves, flat T waves, arrhythmias.
- Treatment: Oral/KCl IV (if severe).

Hypernatremia (Na $^+$ >145) \rightarrow Water loss (diabetes insipidus, dehydration).

Hyponatremia (Na $^+$ <135) \rightarrow SIADH, heart failure, cirrhosis.

3. Acute Kidney Injury (AKI)

Prerenal (\downarrow Renal perfusion)

- Causes: Hypovolemia, HF, NSAIDs, renal artery stenosis.
- Labs: BUN:Cr >20, FENa <1%, urine osmolality >500.
- Treatment: Fluids, correct underlying cause.

Intrinsic Renal

- Acute Tubular Necrosis (ATN):
 - Causes: Ischemia (prolonged hypotension), nephrotoxins (aminoglycosides, contrast).
 - Labs: Muddy brown casts, FENa >2%.
- Glomerulonephritis: Hematuria, proteinuria, RBC casts.
 Interstitial Nephritis: Fever, rash, eosinophilia, WBC casts (drugs: PCN, NSAIDs).

Postrenal (Obstructive)

- Causes: BPH, stones, cervical cancer.
- Diagnosis: Hydronephrosis on ultrasound.
- Treatment: Foley catheter, nephrostomy.

4. Chronic Kidney Disease (CKD)

- Stages: Based on GFR (Stage 5 = ESRD, GFR <15).
- Complications:
 - Anemia: ↓EPO → treat with erythropoietin. Renal osteodystrophy: ↑PTH
 → osteitis fibrosa cystica (treat with vitamin D, phosphate binders).
 - O Metabolic acidosis: Bicarb supplementation.
- Indications for dialysis: Uremia, hyperkalemia, acidosis, fluid overload.

5. Glomerular Diseases

Nephrotic Syndrome (Proteinuria >3.5g/day, hypoalbuminemia, edema, hyperlipidemia)

- Minimal Change Disease: Most common in kids, responds to steroids.
- Focal Segmental Glomerulosclerosis (FSGS): HIV, obesity, heroin use.
- Membranous Nephropathy: Anti-PLA2R antibodies, associated with HBV, malignancy.

Nephritic Syndrome (Hematuria, proteinuria <3.5g, HTN, oliguria)

- Post-Strep GN: 2 weeks after strep throat, subepithelial humps on EM.
- IgA Nephropathy: Hematuria after URI, mesangial IgA deposits.
- Rapidly Progressive GN (RPGN): Crescents on biopsy (Goodpasture's, ANCA vasculitis).

6. Tubular Disorders

- Fanconi Syndrome: Proximal RTA + glycosuria, phosphaturia, aminoaciduria.
- RTA Types:
 - Type 1 (Distal): Hypokalemia, urine pH >5.5 (stones, Sjögren's).
 - Type 2 (Proximal): Hypokalemia, urine pH <5.5 (Fanconi, multiple myeloma).
 - Type 4 (Hypoaldosteronism): Hyperkalemia (DM, ACEIs, spironolactone).

7. Hypertension & Renal Artery Stenosis

- Renovascular HTN: Unilateral small kidney, ↑renin, bruit.
- Diagnosis: Renal artery Doppler, captopril renography.
- Treatment: Revascularization (stenting) or ACEI if bilateral.

8. Dialysis Complications

- Disequilibrium Syndrome: Headache, seizures (rapid solute shifts).
- Access Issues: AV fistula (best), graft, catheter (highest infection risk).

9. High-Yield Pharmacology

- Loop Diuretics (Furosemide): Acts at ascending loop, causes hypokalemia, ototoxicity.
- Thiazides (HCTZ): Acts at DCT, causes hypercalcemia, hypokalemia.
- Spironolactone: K⁺-sparing (blocks aldosterone), used in ascites.

10. Key Formulas

- FENa (%) = (Urine Na × Plasma Cr) / (Plasma Na × Urine Cr) × 100 <1% = Prerenal, >2% = ATN.
- Anion Gap = Na₊ (Cl₋ + HCO_{3_}) (Normal = 8-12).

Here are some high-yield rheumatology facts for the USMLE:

1. Rheumatoid Arthritis (RA)

- Autoantibodies: RF (IgM) and anti-CCP (more specific).
- Pathology: Synovial hyperplasia \rightarrow pannus formation \rightarrow joint destruction.
- Extra-articular: Rheumatoid nodules, interstitial lung disease (ILD), Felty's syndrome (RA + splenomegaly + neutropenia), Sjögren's syndrome (secondary).
- X-ray: Juxta-articular osteopenia, erosions, joint space narrowing.
- Treatment: DMARDs (methotrexate 1st line), TNF- α inhibitors (if refractory).

2. Systemic Lupus Erythematosus (SLE)

- **Diagnosis**: **ANA** (sensitive but not specific), **anti-dsDNA** (specific, correlates with nephritis), **anti-Smith** (highly specific).
- Clinical: Malar rash, discoid rash, photosensitivity, arthritis, serositis (pleuritis/pericarditis), renal disease (lupus nephritis), neuropsychiatric symptoms.
- Drug-induced lupus: Hydralazine, procainamide, isoniazid → anti-histone antibodies.

3. Seronegative Spondyloarthropathies (HLA-B27 associated)

- Ankylosing spondylitis: Sacroiliitis (X-ray: bamboo spine), uveitis, aortic regurgitation.
- Reactive arthritis: Triad (urethritis, conjunctivitis, arthritis) "Can't see, can't pee, can't climb a tree".
- Psoriatic arthritis: DIP joint involvement, nail pitting, "sausage digits" (dactylitis).
- Enteropathic arthritis: Associated with IBD (Crohn's/UC).

4. Gout vs. Pseudogout

- Gout: Monosodium urate crystals (negatively birefringent, needle-shaped).
 - Risk factors: Male, alcohol, diuretics, high-purine diet.
 - Treatment: Acute NSAIDs/colchicine/steroids; Chronic allopurinol/febuxostat (↓ uric acid).
- Pseudogout: Calcium pyrophosphate (CPP) crystals (weakly positively birefringent, rhomboid). O Associated with: Hemochromatosis, hyperparathyroidism, hypomagnesemia.

5. Sjögren's Syndrome

- Autoantibodies: Anti-Ro (SS-A) and Anti-La (SS-B).
- Clinical: Dry eyes (keratoconjunctivitis sicca), dry mouth (xerostomia), parotid enlargement, lymphoma risk.

6. Systemic Sclerosis (Scleroderma)

- Limited (CREST syndrome):
 - o Calcinosis, Raynaud's, Esophageal dysmotility, Sclerodactyly, Telangiectasia.
 - Autoantibody: Anti-centromere.
- Diffuse: Pulmonary fibrosis, renal crisis (malignant HTN + ↑ renin).
 - Autoantibody: Anti-Scl-70 (anti-topoisomerase).

7. Polymyositis (PM) & Dermatomyositis (DM)

- Autoantibodies: Anti-Jo-1 (PM, associated with interstitial lung disease).
- Clinical: Proximal muscle weakness, elevated CK.
- DM-specific: Heliotrope rash, Gottron's papules.
- Associated with malignancy (especially DM).

8. Vasculitides

- Giant Cell (Temporal) Arteritis:
 - Symptoms: Headache, jaw claudication, vision loss (emergency give steroids!).
 - Diagnosis: ESR/CRP ↑, temporal artery biopsy.
- Takayasu's arteritis: "Pulseless disease", aortic arch involvement (young women).
- Granulomatosis with Polyangiitis (GPA, Wegener's):
 - c-ANCA (anti-PR3). O Triad: Necrotizing granulomas (lungs, sinuses), glomerulonephritis.
- Microscopic Polyangiitis (MPA): p-ANCA (anti-MPO), no granulomas.
- Eosinophilic Granulomatosis with Polyangiitis (EGPA, Churg-Strauss): Asthma, eosinophilia, p-ANCA.

9. Osteoarthritis (OA) vs. Rheumatoid Arthritis (RA)

Feature	ОА	RA

Joints	DIP, PIP, knees, hips	MCP, PIP, wrists
Symmetry	Asymmetric early	Symmetric
X-ray	Osteophytes, joint space narrowing	Erosions, juxta-articular osteopenia
Inflammatio n	Minimal	Synovitis, pannus

10. Miscellaneous

- Polymyalgia Rheumatica: Proximal muscle pain/stiffness, ↑ ESR, responds to steroids (often coexists with giant cell arteritis).
- Behçet's Disease: Oral/genital ulcers, uveitis, pathergy.
- Lyme Disease: Erythema migrans, knee arthritis, facial palsy (CN VII).

Key Mnemonics:

- **SLE criteria (SOAP BRAIN MD)** Serositis, Oral ulcers, Arthritis, Photosensitivity, Blood disorders, Renal, ANA, Immunologic, Neurologic, Malar rash, Discoid rash.
- CREST syndrome Calcinosis, Raynaud's, Esophageal dysmotility, Sclerodactyly, Telangiectasia.

RESPIRATORY

1. Obstructive vs. Restrictive Lung Disease

Feature	Obstructive (↓ FEV₁/FVC)	Restrictive (↓ TLC, normal/increased FEV₁/FVC)
Example s	COPD, Asthma, Bronchiectasis	ILD, Sarcoidosis, Kyphoscoliosis, Obesity
Spiromet ry	FEV ₁ /FVC < 0.7	FEV₁/FVC ≥ 0.7, ↓ TLC
Causes	Airway obstruction	Reduced lung expansion (fibrosis, chest wall)

2. Chronic Obstructive Pulmonary Disease (COPD)

- Types:
 - Chronic bronchitis ("Blue bloater") productive cough, hypoxemia, cor pulmonale. Emphysema ("Pink puffer") dyspnea, barrel chest, ↓ DLCO, α-1 antitrypsin deficiency (panacinar emphysema in young non-smokers).
- Pathology: Destruction of alveoli (emphysema) or mucus hypersecretion (bronchitis).
- Diagnosis: Post-bronchodilator FEV₁/FVC < 0.7.
- Treatment: Smoking cessation, SABA/LAMA (bronchodilators), oxygen if hypoxemic.

3. Asthma

- Triad: Wheezing, dyspnea, cough (worse at night).
- Pathology: Reversible airway obstruction, eosinophilic inflammation.
- Diagnosis: ↑ FEV₁ after bronchodilator (≥12%), ↓ FEV₁/FVC.
- Severe attack: Silent chest, PaCO₂ > 40 mmHg (ominous sign of respiratory fatigue).
- Treatment: SABA (albuterol), ICS, leukotriene inhibitors (montelukast).

4. Pulmonary Embolism (PE)

- Wells Criteria (estimates probability):
 - Clinical signs of DVT, HR > 100, recent immobilization/surgery, hemoptysis, malignancy.
- Diagnosis: CT angiography (gold standard), D-dimer (if low probability).

- ECG: S1Q3T3, right heart strain (RBBB, RAD, T-wave inversions V1-V4).
- Treatment: Anticoagulation (heparin → DOACs/warfarin).

5. Pneumonia

Туре	Key Features	Causative Organisms
Communit y (CAP)	Fever, cough, crackles	S. pneumoniae (most common), H. influenzae, Mycoplasma (atypical)
Hospital (HAP)	>48h after admission	Pseudomonas, MRSA, Klebsiella
Aspiration	Right lower lobe, foulsmelling sputum	Anaerobes (Bacteroides, Peptostreptococcus)
тв	Night sweats, weight loss, +AFB stain	Mycobacterium tuberculosis

- Diagnosis: CXR, sputum culture, procalcitonin (bacterial vs. viral).
- Treatment: CAP Ceftriaxone + azithromycin; HAP Piperacillintazobactam/vancomycin.

6. Interstitial Lung Disease (ILD)

- Causes: Pneumoconioses (asbestosis, silicosis), drugs (amiodarone, bleomycin), sarcoidosis, idiopathic pulmonary fibrosis (IPF).
- Findings: Bibasilar crackles, restrictive pattern, honeycombing on CT.
- Diagnosis: HRCT, lung biopsy.

7. Lung Cancer

- Non-small cell (NSCLC) (80%):
 - o Adenocarcinoma (peripheral, non-smokers, EGFR mutations).
 - Squamous cell (central, smoking, PTHrP → hypercalcemia).
- Small cell (SCLC) (20%): Central, smoking, paraneoplastic syndromes (SIADH, Lambert-Eaton).
- Diagnosis: CT-guided biopsy, PET scan for staging.

8. Pleural Diseases

- Pleural effusion:
 - Transudate (↑ hydrostatic pressure CHF, nephrotic syndrome).
 - Exudate (↑ permeability pneumonia, malignancy, TB) Light's criteria.
- Pneumothorax: O Primary (tall, thin males, ruptured blebs). O Secondary (COPD, trauma).
 - Tension pneumo (tracheal deviation, hypotension, absent breath sounds).

9. Sleep Apnea

- Obstructive (OSA): Daytime sleepiness, snoring, obesity, hypertension.
- Diagnosis: Polysomnography (PSG).
- Treatment: CPAP, weight loss.

10. High-Altitude Pulmonary Edema (HAPE)

- Symptoms: Dyspnea, cough, pink frothy sputum.
- Pathophysiology: Hypoxia → pulmonary vasoconstriction → edema.
- Treatment: Descent, oxygen, nifedipine.

Key Mnemonics:

- COPD exacerbation treatment: Steroids, Oxygen, Bronchodilators, Antibiotics (if infective).
- Asthma drugs (quick relief vs. maintenance): SABA (rescue), ICS (prevention).
- Pneumonia bugs: CAP = S. pneumo, HAP = Pseudomonas/MRSA, Atypical = Mycoplasma/Chlamydia

Final Tip: Focus on clinical correlations (e.g., nerve injuries, hernias, fractures) and vascular supply patterns for exams!

Source: Dr. Maurych's *Anatomy Shelf Notes*.

High-yield physiology concepts from Guyton & Hall that are essential for USMLE/NRE:

1. Cardiovascular Physiology

- Frank-Starling Law: ↑ Preload → ↑ Stroke volume (up to a point)
- Cardiac Output (CO) = HR × SV O SV depends on preload, afterload, contractility
- Vascular Resistance:
 - o **Poiseuille's Law**: Resistance ∝ 1/radius⁴ (most important regulator of blood flow)
- Coronary Blood Flow:
 - Occurs mainly during diastole (LV compression blocks flow in systole)

2. Renal Physiology

- GFR = Kf × (PGC PBS π GC) \odot Autoregulation: Myogenic + Tubuloglomerular feedback (macula densa)
- Na+ Reabsorption:
 - PCT (67%) Na+/H+ exchanger, Loop of Henle (25%) NKCC2, DCT (5%) Na+/Cl– cotransporter
- Countercurrent Mechanism:
 - Descending limb: Water reabsorption (via aquaporins) Ascending limb: Na+/K+/Cl– reabsorption (dilutes urine)
- ADH (Vasopressin):
 \(\backslash\) Water reabsorption in collecting duct (via aquaporin-2)
- Aldosterone: ↑ Na+ reabsorption & K+ secretion in DCT/CD

3. Respiratory Physiology

- Alveolar Gas Equation: PAO₂ = FiO₂(PATM PH₂O) (PaCO₂/R) A-a Gradient = PAO₂ PaO₂ (↑ in V/Q mismatch, diffusion defects, R→L shunt)
- Oxygen-Hemoglobin Dissociation Curve:
 - Right shift (↓ affinity): ↑ CO₂, ↑ H₊, ↑ 2,3-BPG, ↑ Temp (e.g., exercise) Left shift (↑ affinity): ↓ CO₂, ↓ H₊, ↓ 2,3-BPG, ↓ Temp (e.g., fetal Hb)
- Ventilation-Perfusion (V/Q) Mismatch: Dead space (V/Q = ∞): Pulmonary embolism
 Shunt (V/Q = 0): Atelectasis, pneumonia

4. Acid-Base Balance

- Henderson-Hasselbalch Equation: pH = 6.1 + log([HCO_{3_}] / (0.03 × PCO₂))
- Metabolic Acidosis: ↓ HCO₃ (e.g., DKA, diarrhea) → ↑ Anion Gap (MUDPILES)
 Respiratory Acidosis: ↑ PCO₂ (e.g., COPD, opioid OD)
 Renal Compensation:
 - Proximal tubule: Reabsorbs HCO₃_
 - Collecting duct: Secretes H₊ (via H₊-ATPase)

5. Neurophysiology

- Action Potential:
 - Depolarization: Na+ influx (fast channels) Repolarization: K+ efflux (slow channels)
 - Refractory Periods: Absolute (Na+ inactivation) vs. Relative (K+ efflux)
- Synaptic Transmission:
 - EPSP (Na+/Ca² influx) vs. IPSP (Cl_influx/K efflux)
 - o **NMJ**: ACh → Nicotinic receptors → Na+ influx → Muscle contraction

6. Endocrine Physiology

- Hypothalamic-Pituitary Axis:
 - TRH \rightarrow TSH \rightarrow T3/T4 (\uparrow BMR, \uparrow β -receptors)
 - CRH → ACTH → Cortisol (↑ gluconeogenesis, immunosuppression)
- Insulin vs. Glucagon:
 - o Insulin (β-cells): \uparrow Glucose uptake (GLUT4), \downarrow Lipolysis o Glucagon (α -cells): \uparrow Glycogenolysis, \uparrow Gluconeogenesis

7. GI Physiology

- Gastric Acid Secretion:
 - Parietal cells: H₊/K₊ ATPase (stimulated by ACh, gastrin, histamine)
 Cephalic phase: Vagal stimulation (ACh)
- Bile: Emulsifies fats (bile salts), excretes bilirubin (no digestion)
- Absorption Sites:

- o Iron/Calcium: Duodenum
- o B12/BILE SALTS: Ileum

8. Hematology

- · Hemoglobin:
 - O_2 binding: Cooperative binding (sigmoidal curve) CO Poisoning: \uparrow COHb \rightarrow Left shift (\downarrow O_2 delivery)
- Clotting Cascade:
 - Extrinsic Pathway: Tissue factor (Factor VIIa) \circ Intrinsic Pathway: XII \rightarrow XI \rightarrow IX \rightarrow X

9. Autonomic Nervous System

- Sympathetic (Fight/Flight):
 - α_1 : Vasoconstriction (\uparrow BP) β_1 : \uparrow HR, \uparrow Contractility
- Parasympathetic (Rest/Digest):
 - M3: ↑ Secretions, ↓ HR (vagal tone)

10. Special Senses

- Vision:
 - o Rods: Night vision (rhodopsin) o Cones: Color vision (S/M/L opsins)
- Hearing:
 - o **Organ of Corti**: Hair cells (stereocilia bend → K+ influx)

Key Mnemonics for USMLE

- Anion Gap Metabolic Acidosis: MUDPILES (Methanol, Uremia, DKA, Paraldehyde, INH, Lactic acidosis, Ethylene glycol, Salicylates)
- Cushing's Triad: ↑ BP, ↓ HR, Irregular RR (sign of ↑ ICP)
- Left Shift (O₂-Hb Curve): CADET face right (CO, Acidosis, 2,3-DPG, Exercise, Temperature)

Final Tips

- Master these equations: Nernst, Fick's Law, Henderson-Hasselbalch.
- Focus on clinical correlations: e.g., V/Q mismatch (PE vs. pneumonia), acidbase disorders, autonomic drug effects.

High-Yield <u>Biochemistry</u> Points from <u>Lippincott Illustrated Reviews</u> (USMLE Focus)

1. Enzyme Kinetics & Regulation

- Michaelis-Menten Equation:
 - $V_0 = (V_{max} \times [S]) / (K_m + [S])$ $K_m = Substrate$ concentration at $\frac{1}{2}V_{max}$ ($\frac{1}{2}V_{max}$).
- Inhibitors:
 - **Competitive**: \uparrow K_m (e.g., statins for HMG-CoA reductase). **Noncompetitive**: \downarrow V_{max} (e.g., cyanide for cytochrome oxidase).
 - **Uncompetitive**: \downarrow both K_m and V_{max} .
- Allosteric Regulation: Feedback inhibition (e.g., CTP inhibits aspartate transcarbamoylase).

2. Carbohydrate Metabolism

- Glycolysis:
 - Rate-limiting enzyme: Phosphofructokinase-1 (PFK-1) (activated by AMP/F2,6BP; inhibited by ATP/citrate).
 - o **ATP Yield**: 2 ATP (anaerobic), 30-32 ATP (aerobic).
- Gluconeogenesis:
 - Key enzymes: Pyruvate carboxylase, PEP carboxykinase, F1,6bisphosphatase, G6Pase
 - o **Fed state**: Insulin \uparrow glycolysis, \downarrow gluconeogenesis.
- TCA Cycle:
 - Rate-limiting enzyme: Isocitrate dehydrogenase (NAD₊ \rightarrow NADH). ATP Yield: 3 NADH, 1 FADH₂, 1 GTP per cycle.

3. Lipid Metabolism

- Fatty Acid Oxidation:
 - o **β-Oxidation**: Occurs in mitochondria; **Carnitine shuttle** transports FA.
 - **Deficiency**: MCAD deficiency \rightarrow hypoglycemia, \uparrow dicarboxylic acids.
- Ketogenesis:
 - Liver mitochondria (starvation/DKA); HMG-CoA synthase is key.
- Cholesterol Synthesis:
 - o **HMG-CoA reductase** (rate-limiting; inhibited by statins).

4. Nitrogen Metabolism

- Urea Cycle:
 - Location: Liver (mitochondria + cytoplasm). Rate-limiting enzyme: Carbamoyl phosphate synthetase I (requires Nacetylglutamate). Deficiency: Ornithine transcarbamylase (OTC) → ↑ ammonia, ↑ orotic acid (X-linked).
- Amino Acid Catabolism:
 - Branched-chain AAs (Leu, Ile, Val): Defects cause maple syrup urine disease.
 Phenylalanine: PKU → ↑ phenylalanine, ↓ tyrosine (treated with low-Phe diet).

5. Vitamins & Cofactors

- B1 (Thiamine):
 - Deficiency: Beriberi (wet/dry), Wernicke-Korsakoff.
 - o **Enzymes**: PDH, α -KGDH, transketolase.
- B12 (Cobalamin):
 - Deficiency: Megaloblastic anemia, neurological defects (↓ methionine synthase).
- Folate:
 - Deficiency: Neural tube defects, megaloblastic anemia (↓ dTMP synthesis).

6. Molecular Biology

- DNA Replication:
 - **Leading strand**: Continuous $(5' \rightarrow 3')$.
 - o Lagging strand: Okazaki fragments (RNA primer needed).
- Mutations:
 - **Missense**: Single AA change (e.g., sickle cell \rightarrow Glu \rightarrow Val). **Nonsense**: Premature stop codon (e.g., Duchenne muscular dystrophy).

7. Glycogen Storage Diseases

Disease	Deficient Enzyme	Key Feature
Von Gierke	G6Pase	Hepatomegaly, hypoglycemia
McArdle	Muscle glycogen phosphorylase	Exercise intolerance, myoglobinuria
Pompe	Lysosomal α-1,4-glucosidase	Cardiomegaly, infantile death

8. Metabolic Pathways Integration

- **Fed State**: **Insulin** ↑ glycolysis, FA synthesis, glycogenesis.
- Fasting State: Glucagon ↑ gluconeogenesis, glycogenolysis, ketogenesis.

9. Clinical Correlations

- G6PD Deficiency: Hemolytic anemia with oxidative stress (favism, sulfa drugs).
- **Lesch-Nyhan Syndrome**: HGPRT deficiency → ↑ uric acid, self-mutilation.
- **Alkaptonuria**: Homogentisate oxidase deficiency → black urine, ochronosis.

10. High-Yield Mnemonics

- Rate-limiting enzymes:
 - Glycolysis: PFK-1 (People Fear Keto).
 - TCA: Isocitrate DH (I Like TCA).
- Urea Cycle Defects: OTC (Ornithine = Only Treatable with Carbs).

Final Tips for USMLE

- **Focus on diseases** (e.g., glycogen storage, vitamin deficiencies).
- Master rate-limiting enzymes and their regulators.
- Link pathways to clinical scenarios (e.g., DKA → ketogenesis, PKU → tyrosine deficiency).

High-Yield General Pathology Points from Robbins Pathology (USMLE Focus)

1. Cellular Adaptations & Injury

- Atrophy: ↓ Cell size (e.g., disuse atrophy, denervation).
- **Hypertrophy**: ↑ Cell size (e.g., cardiac muscle in hypertension).
- **Hyperplasia**: ↑ Cell number (e.g., endometrial hyperplasia, BPH).
- **Metaplasia**: Replacement with another cell type (e.g., Barrett's esophagus, squamous metaplasia in smokers).
- Reversible Injury: Cellular swelling, fatty change.
- Irreversible Injury: Necrosis (coagulative, liquefactive, caseous, fat, fibrinoid) vs. Apoptosis (programmed cell death).

2. Inflammation & Repair

Acute Inflammation:

- Cardinal signs: Rubor (redness), tumor (swelling), calor (heat), dolor (pain), functio laesa (loss of function).
- Mediators: Histamine (mast cells), prostaglandins (COX pathway), bradykinin (pain).
- Outcomes: Resolution, abscess, fibrosis, chronic inflammation.

Chronic Inflammation:

- Cells: Macrophages, lymphocytes, plasma cells.
- o **Granuloma**: Epithelioid macrophages + giant cells (e.g., TB, sarcoidosis).

Wound Healing:

- Primary intention: Clean surgical incision.
- Secondary intention: Large wound (e.g., ulcer).

3. Hemodynamic Disorders

- **Edema**: ↑ Hydrostatic pressure (e.g., CHF) or ↓ oncotic pressure (e.g., nephrotic syndrome).
- Thrombosis: Virchow's triad (endothelial injury, stasis, hypercoagulability).
- Embolism:
 - **Pulmonary embolism**: DVT \rightarrow right heart \rightarrow lungs.
 - Paradoxical embolism: DVT → ASD/VSD → systemic circulation.

Infarction:

Pale infarct: Arterial occlusion (e.g., kidney, heart). O Hemorrhagic infarct:
 Venous occlusion or dual blood supply (e.g., lung, liver).

4. Neoplasia

- Benign vs. Malignant:
 - o **Benign**: Well-differentiated, slow growth, no metastasis.
 - o Malignant: Poorly differentiated, invasion/metastasis.
- Carcinogenesis:
 - o **Oncogenes**: RAS (most common), MYC, HER2/neu.
 - o **Tumor suppressors**: p53 ("guardian of genome"), RB, APC.
- Paraneoplastic Syndromes:
 - o **Small cell lung cancer**: ACTH (Cushing's), SIADH.
 - Squamous cell carcinoma: PTHrP (hypercalcemia).

5. Genetic Disorders

- Autosomal Dominant:
 - o **Huntington's disease**: CAG repeats → chorea, dementia.
 - o Familial adenomatous polyposis (FAP): APC mutation \rightarrow colon polyps.
- Autosomal Recessive:
 - **Cystic fibrosis**: Δ F508 mutation in *CFTR* \rightarrow \uparrow Cl_ in sweat, lung/pancreas disease.
 - o Phenylketonuria (PKU): Phenylalanine hydroxylase deficiency.
- X-Linked:
 - \circ **Duchenne muscular dystrophy**: Dystrophin deletion $\rightarrow \uparrow$ CK, Gower's sign.

6. Immunopathology

- Hypersensitivity Reactions:
 - Type I (IgE): Anaphylaxis, asthma. Type II (IgG/IgM): AIHA, Goodpasture's (anti-GBM). ○ Type III (Immune complexes): SLE, post-streptococcal GN.
 - Type IV (T-cell): TB granuloma, contact dermatitis.
- Autoimmune Diseases:
 - SLE: Anti-dsDNA, anti-Smith, butterfly rash.
 - o **Rheumatoid arthritis**: Anti-CCP, synovial pannus.

7. Infectious Disease Pathology

- Bacterial:
 - Staph. aureus: Abscesses (coagulase +), toxic shock syndrome (TSST-1).
 - o **Strep. pyogenes**: Rheumatic fever (M protein), post-streptococcal GN.
- Viral:
 - **HIV**: \downarrow CD4₊ T cells \rightarrow opportunistic infections (PCP, CMV, TB).
 - HPV: Squamous cell carcinoma (types 16/18).

8. Environmental & Nutritional Pathology

- **Smoking**: Squamous metaplasia \rightarrow lung cancer, emphysema (α 1-antitrypsin deficiency).
- Alcohol: Fatty liver → cirrhosis, Wernicke-Korsakoff (B1 deficiency).
- Vitamin Deficiencies:
 - o **B1 (Thiamine)**: Wet/dry beriberi, Wernicke-Korsakoff.
 - o **B12**: Megaloblastic anemia, subacute combined degeneration.

9. High-Yield Mnemonics

- Necrosis Types: CLF (Coagulative, Liquefactive, Fat).
- Tumor Suppressors: p53, RB, APC ("PRAy they don't mutate").
- Granulomatous Diseases: TB, Sarcoidosis, Cat-scratch, Histoplasmosis (TSCH).

Final Tips for USMLE

- **Focus on mechanisms**: e.g., how *p53* mutations cause Li-Fraumeni syndrome.
- Link pathology to clinical findings: e.g., caseous necrosis → TB.
- Master high-yield images: e.g., Reed-Sternberg cells (Hodgkin's lymphoma).

High-Yield Community Medicine Points for USMLE

1. Epidemiology & Biostatistics

- Prevalence vs. Incidence:
 - Prevalence = Total cases / Total population at a given time.
 - Incidence = New cases / Population at risk over a time period.
- Study Types:
 - o Randomized Controlled Trial (RCT) = Gold standard for causation.
 - Cohort Study = Prospective, calculates relative risk (RR).
 - Case-Control Study = Retrospective, calculates odds ratio (OR).
- Sensitivity & Specificity:
 - Sensitivity = True positives / (True positives + False negatives).
 - Specificity = True negatives / (True negatives + False positives).

2. Screening Tests

- Positive Predictive Value (PPV) = True positives / (True positives + False positives).
- Negative Predictive Value (NPV) = True negatives / (True negatives + False negatives).
- High Sensitivity Tests = Best for ruling out disease (SnOUT).
- High Specificity Tests = Best for ruling in disease (SpIN).

3. Vaccination & Immunization

- Live Attenuated Vaccines (Contraindicated in pregnancy/immunocompromised):
 - o MMR, Varicella, Yellow Fever, Rotavirus, Oral Polio (Sabin).
- Inactivated/Killed Vaccines:
 - o Influenza, Hepatitis A, Rabies, Salk Polio.
- Toxoid Vaccines:
 - o Tetanus, Diphtheria.

4. Infectious Disease Control

- Quarantine = Separates exposed but asymptomatic individuals.
- Isolation = Separates symptomatic infected individuals.
- R0 (Basic Reproduction Number): R0 > 1 = Epidemic likely.
 - o R0 < 1 = Outbreak will die out.

5. Occupational & Environmental Health

- Lead Poisoning:
 - Symptoms: Anemia, wrist drop, encephalopathy.

- o **Treatment**: Chelation (EDTA, Succimer).
- Asbestos Exposure: O Diseases: Mesothelioma, lung cancer, asbestosis.

6. Maternal & Child Health

- APGAR Score (Assessed at 1 & 5 min):
 O-3 = Severe distress.
 O-4-6 = Moderate distress.
 - o 7-10 = Normal.
- Breastfeeding Benefits: O Colostrum = Rich in IgA, protects against infections.

7. Nutrition & Deficiency Disorders

- Vitamin A Deficiency: O Symptoms: Night blindness, Bitot's spots, xerophthalmia.
- Vitamin D Deficiency: O Children: Rickets (bowing of legs). O Adults: Osteomalacia.

8. Preventive Medicine

- Primary Prevention = Prevents disease (e.g., vaccination).
- **Secondary Prevention** = Early detection (e.g., screening).
- Tertiary Prevention = Reduces complications (e.g., rehab).

9. High-Yield Mnemonics

- Live Vaccines: MY VRR (MMR, Varicella, Rotavirus, Rubella, Yellow Fever).
- Notifiable Diseases: Mumps, Measles, Rabies, TB, Hepatitis A/B/C.

Final Tips for USMLE

- Focus on screening tests, vaccination schedules, and R0 calculations.
- Memorize key deficiency diseases and their treatments.
- Link epidemiology to public health interventions.

Or Abdull Basit Luloali